

Refine Search

Search Results -

Terms	Documents
6805879.pn.	1

Database:

US Pre-Grant Publication Full-Text Database
US Patents Full-Text Database
US OCR Full-Text Database
EPO Abstracts Database
JPO Abstracts Database
Derwent World Patents Index
IBM Technical Disclosure Bulletins

Search:

L3

[Refine Search](#)[Recall Text](#)[Clear](#)[Interrupt](#)

Search History

DATE: Sunday, September 10, 2006 [Purge Queries](#) [Printable Copy](#) [Create Case](#)

<u>Set Name</u> side by side	<u>Query</u>	<u>Hit Count</u>	<u>Set Name</u> result set
<i>DB=USPT; PLUR=YES; OP=OR</i>			
<u>L3</u>	6805879.pn.	1	<u>L3</u>
<u>L2</u>	6284375.pn.	1	<u>L2</u>
<i>DB=PGPB,USPT; PLUR=YES; OP=OR</i>			
<u>L1</u>	Tuo near Jin	13	<u>L1</u>

END OF SEARCH HISTORY

Refine Search

Your wildcard search against 10000 terms has yielded the results below.

Your result set for the last L# is incomplete.

The probable cause is use of unlimited truncation. Revise your search strategy to use limited truncation.

Search Results -

Terms	Documents
L18 and (por\$ same (alumina or silica or cellulose))	14

Database:

US Pre-Grant Publication Full-Text Database
US Patents Full-Text Database
US OCR Full-Text Database
EPO Abstracts Database
JPO Abstracts Database
Derwent World Patents Index
IBM Technical Disclosure Bulletins

Search:

L19

Refine Search

Recall Text

Clear

Interrupt

Search History

DATE: Sunday, September 10, 2006 [Purge Queries](#) [Printable Copy](#) [Create Case](#)

<u>Set</u> <u>Name</u>	<u>Query</u>	<u>Hit</u> <u>Count</u>	<u>Set</u> <u>Name</u> result set
	DB=PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD; PLUR=YES; OP=OR		
<u>L19</u>	L18 and (por\$ same (alumina or silica or cellulose))	14	<u>L19</u>
<u>L18</u>	L17 and (capsule or tablet or granule or (coated adj (tablet or granule)))	1116	<u>L18</u>
<u>L17</u>	L15 and carrier	1121	<u>L17</u>
<u>L16</u>	L15 and (por\$ same (alumina or silica or cellulose))	15	<u>L16</u>
<u>L15</u>	L8 and (alumina or silica or cellulose)	1136	<u>L15</u>
<u>L14</u>	L12 and (por\$ same (alumina or silica or cellulose))	2	<u>L14</u>
<u>L13</u>	L12 and (por\$ near5 (alumina or silica or cellulose))	1	<u>L13</u>
<u>L12</u>	L11 and (alumina or silica or cellulose)	117	<u>L12</u>
<u>L11</u>	L8 and (cyclosporine or triamterene or acyclovir or doxorubicin or labetalol or doxepin or methyl dopa or pentoxifill)	129	<u>L11</u>

<u>L10</u>	L8 and (cyclosporine or triamterene or acyclovir, or doxorubicin or labetalol or doxepin or methyldopa or pentoxifill)	129	<u>L10</u>
<u>L9</u>	L8 and (surface adj area)	61	<u>L9</u>
<u>L8</u>	L5 and emulsion	1220	<u>L8</u>
<u>L7</u>	L5 and (porous near5 powder)	28	<u>L7</u>
<u>L6</u>	L5 and (porous adj powder)	4	<u>L6</u>
<u>L5</u>	L4 and (powder adj solution)	1308	<u>L5</u>
<u>L4</u>	L3 and (powder same (lipid or phospholipid or surfactant))	11595	<u>L4</u>
<u>L3</u>	(porous or tablet\$ or (compress\$ or (free near flowing))) near10 powder	152833	<u>L3</u>
<i>DB=USPT; PLUR=YES; OP=OR</i>			
<u>L2</u>	6241997.pn.	1	<u>L2</u>
<u>L1</u>	6280770.pn.	1	<u>L1</u>

END OF SEARCH HISTORY

(FILE 'HOME' ENTERED AT 19:51:48 ON 10 SEP 2006)

FILE 'CAPLUS, MEDLINE, USPATFULL' ENTERED AT 19:52:08 ON 10 SEP 2006

L1	1838 S POWDER(W)SOLUTION
L2	32 S L1 AND (POR? (5A) (ALUMINA OR SILICA OR CELLULOSE))
L3	32 DUPLICATE REMOVE L2 (0 DUPLICATES REMOVED)
L4	32 FOCUS L3 1-
L5	38 S L1 (P) (COMPRESS? OR (FREE(W)FLOWING) OR POROUS)
L6	3 S L5 AND (DISOL? OR DISSOLUTION OR DISPERS?) (P) (INSOLUBLE OR
L7	3 DUPLICATE REMOVE L6 (0 DUPLICATES REMOVED)
L8	0 S L1 (P) (INSOLUBLE (5A) (DRUG OR PHARMACEUTICAL))
L9	374 S L1 AND POROUS
L10	215 S L9 AND (INSOLUBLE (P) (DRUG OR MEDICAMENT OR PHARMACEUTICAL)
L11	1 S L10 AND (POROUS (10A) (ALUMINA OR SILICA OR CELLULOSE))

L11 ANSWER 1 OF 1 USPATFULL on STN

TI Solid dosage forms for rapid dissolution of poorly soluble drugs

AB This invention demonstrated novel pharmaceutical compositions that improve dissolution, water dispersion and/or oral absorption of insoluble or poorly soluble drugs without increase in formulation complicity and patient appliance as compared with conventional solid-dosage form. The compositions of the present invention comprise a lipid or mixed lipids that dissolve the insoluble or poorly soluble drugs and forms solution, micelles, microemulsion or emulsion with the drugs in aqueous media. The compositions further comprise a porous powder or mixed porous powder that absorb the drug-lipid melts in a considerable amount (>than their own mass) while remaining free flowing and compressible in nature. Due to their excellent effectiveness-simplicity ratio, the compositions of this invention have a wide applicability to therapeutic compounds whose efficacy is limited by poor solubility, low dissolution rate and less absorption.

ACCESSION NUMBER: 2004:1869 USPATFULL

TITLE: Solid dosage forms for rapid dissolution of poorly soluble drugs

INVENTOR(S): Jin, Tuo, Highland Park, NJ, UNITED STATES

PATENT ASSIGNEE(S): BioPharm Solutions Inc. (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2004001888	A1	20040101
APPLICATION INFO.:	US 2003-606344	A1	20030626 (10)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2002-391756P	20020626 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	Albert Wai-Kit Chan, Law Offices of Albert Wai-Kit Chan, LLC, World Plaza, Suite 604, 141-07 20th Avenue, Whitestone, NY, 11357	
NUMBER OF CLAIMS:	19	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	5 Drawing Page(s)	
LINE COUNT:	632	

CAS INDEXING IS AVAILABLE FOR THIS PATENT